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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO	
09/779,447 02/09/2001		Dipak K. Banerjee	P19850.p06	6690	
7055	7590 05/17/2004		EXAMINER		
	UM & BERNSTEIN, I	OWENS JR, HOWARD V			
1950 ROLAI RESTON, V	ND CLARKE PLACE /A 20191		ART UNIT	PAPER NUMBER	
1			1623		
			DATE MAILED: 05/17/200	4	

Please find below and/or attached an Office communication concerning this application or proceeding.

		Applicati	on N	Applicant(s)				
Office Action Summary		09/779,4		BANERJEE ET AL.				
		Examine		Art Unit				
		Howard V		1623				
-	- The MAILING DATE of this communication			,	dress			
Period for Reply								
THE N - Extens after S - If the p - If NO - Failure Any re	PRTENED STATUTORY PERIOD FOR REMAILING DATE OF THIS COMMUNICATION is sions of time may be available under the provisions of 37 CF SIX (6) MONTHS from the mailing date of this communication period for reply specified above is less than thirty (30) days, is period for reply is specified above, the maximum statutory properly within the set or extended period for reply will, by supply received by the Office later than three months after the right days are the remaining and patent term adjustment. See 37 CFR 1.704(b).	DN. FR 1.136(a). In no ev n. a reply within the sta eriod will apply and w statute, cause the app	ent, however, may a reply be tim tutory minimum of thirty (30) days rill expire SIX (6) MONTHS from blication to become ABANDONE	nely filed s will be considered timely the mailing date of this co O (35 U.S.C. § 133).				
Status								
1)🛛	Responsive to communication(s) filed on 1	12 March 2004						
·	This action is FINAL . 2b) This action is non-final.							
3)□	Since this application is in condition for allowance except for formal matters, prosecution as to the merits is							
(closed in accordance with the practice under Ex parte Quayle, 1935 C.D. 11, 453 O.G. 213.							
Dispositio	on of Claims							
4)🛛	Claim(s) <u>2-18</u> is/are pending in the applica	ition.						
•	4a) Of the above claim(s) is/are withdrawn from consideration.							
	5) Claim(s) is/are allowed.							
6)🖂	⊠ Claim(s) <u>2-18</u> is/are rejected.							
7) 🗌	Claim(s) is/are objected to.							
8) 🗌 (Claim(s) are subject to restriction a	nd/or election r	equirement.					
Application	on Papers							
9)□ T	he specification is objected to by the Exar	miner.						
10)⊠ The drawing(s) filed on <u>9/12/2001</u> is/are: a)□ accepted or b)□ objected to by the Examiner.								
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).								
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).								
11) The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.								
Priority u	nder 35 U.S.C. § 119							
a)[Acknowledgment is made of a claim for ford All b) Some * c) None of: 1. Certified copies of the priority docum			-(d) or (f).				
:	Certified copies of the priority docum	nents have bee	en received in Application	on No				
;	3. Copies of the certified copies of the	•		d in this National	Stage			
* 0.	application from the International Bu	•	` ''	ي.				
30	ee the attached detailed Office action for a	riist of the certi	ned copies not receive	a.				
Attachment((s)							
1) Notice	of References Cited (PTO-892)		4) Interview Summary					
2) Notice of Draftsperson's Patent Drawing Review (PTO-948) Paper No(s)/Mail Date 3) Information Disclosure Statement(s) (PTO-1449 or PTO/SB/08) 5) Notice of Informal Patent Application (PTO-152)								
3) Information Disclosure Statement(s) (PTO-1449 or PTO/SB/08) Paper No(s)/Mail Date 5) Notice of Informal Patent Application (PTO-152) 6) Other:								

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Response to Arguments

The following is in response to the amendment filed 3/12/04:

An action on the merits of claims 2-18 is contained herein below.

The text of those sections of Title 35, U.S. Code not included in this action can be found in a prior Office action.

Finality of the last office action is withdrawn.

Claim for Domestic Priority

The claim for domestic priority under 35 U.S.C. 119(e) to provisional application no. 60/181,312 is acknowledged.

Request for Approval of Drawing Corrections

The drawings were received on September 12, 2001. These drawings are approved.

Response to Restriction Requirement

Applicant's election with traverse of Group I in Paper No. 11 is acknowledged. It is noted that Groups I-III have been acted on by the examiner and thus comments regarding the examination of these groups for restriction purposes are moot. The traversal for groups IV - XIV is on the ground(s) that an appropriate explanation of how a serious burden would be imposed has not been stated by the examiner. This is not found persuasive because as cited in the restriction requirement mailed 1/6/03, "these inventions are distinct for the reasons given above and have acquired a separate status in the art because of their recognized divergent subject matter, restriction for examination purposes as indicated is proper. To search fourteen distinct and divergent inventions would impose an undue burden upon an examiner assigned this application". Thus an explanation of why an undue burden would be imposed was set forth;

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moreover, applicant has not responded as to how the examiner was in err with respect to the distinct and divergent classification of the inventions set forth in Groups IV-XIV.

This application contains claims drawn to an invention nonelected with traverse in Paper No. 11. A complete reply to the final rejection must include cancellation of nonelected claims or other appropriate action (37 CFR 1.144) See MPEP § 821.01. The requirement is still deemed proper and is therefore made **FINAL**.

35 U.S.C. 103

Applicant's arguments filed 3/12/04 have been fully considered but they are not persuasive.

The rejection of claims 2-18 under 35 U.S.C. 103 as being unpatentable over Banerjee et al., *Indian J. Biochem. and Biophysics*, vol. 30(6), pp. 389-94 and Tiganis et al., *Exp. Cell Research*, vol. 198, pp. 191-200 (1992) is maintained for the reasons of record..

Claims 2-14 and 16-18 are drawn to a method for inhibiting angiogenesis comprising administering a pyrimidine nucleoside, wherein the nucleoside comprises N-acetylated glucosamine or comprises tunicamycin and functional derivatives thereof.

Claim 15 is drawn to the method of claim 1, wherein the inhibition of angiogenesis occurs in a variety of diseases wherein inhibition of angiogenesis would be beneficial, i.e. diabetic retinopathy, artherosclerotic plagues, schleroderma, etc.

Banerjee teaches that angiogenesis comprises (1)endothelial cell proliferation and (2) differentiation into blood capillaries. Banerjee teaches the use of a pyrimidine nucleoside as an antiangiogenic agent as it teaches that the angiogenic process of capillary endothelial cell proliferation is linked to the synthesis of N-linked oligosaccharide chains which is inhibited by the pyrimidine nucleoside tunicamycin (which contains a linked glucosamine). Tiganis et al., further supports the recognition in the prior art of the inhibition of N-glycosylation by tunicamycin and the disruption of vascular proliferation or angiogenesis. Tiganis teaches that the inhibition of glycoproteins by tunicamycin impairs the cell adhesion and the functional properties of

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X,

the endothelial lining of the blood vessels. Thus one of skill in the art would have a reasonable expectation of success that if Tunicamycin is a potent inhibitor of N-glycosylation and that this inhibition disrupts component (1) of angiogenesis, there is clearly a reasonable expectation of success in the use of tunicamycin as an agent which would inhibit angiogenesis.

Aside from presenting new arguments, applicant should note that there has been no specific response by applicant to the discussion of homologs set forth in the prior office action with regards to the Banerjee reference. As cited in the specific office action Banerjee does not specifically mention the claimed homologs of tunicamycin nor the various diseases wherein angiogenesis may be present; however, characteristics normally possessed by members of a homologous series are principally the same, chemists would in general know what to expect in adjacent members of homologs of a known compound. The test of patentability of a compound that is a homologue of a prior art compound is whether the claimed compound possesses beneficial characteristics which are unexpected and unobvious. One of skill in the art would have a reasonable expectation of success in the use of homologs of tunicamycin as angiogenic compounds given the efficacy of the parent compound. There is not data in the prior art nor the specification that presents some property of these homologs apart from that of the parent compound, chiefly the inhibition of angiogenesis. One of skill in the art would also have a reasonable expectation of success that a compound which inhibits angiogenesis would be beneficial in various disease states which may be disrupted by or thrive on the process of angiogenesis. Applicant's claims regarding the administration timetable of the know compound is not patentable given that one of skill in the art practicing the administration of any medical compound determines the optimum dosage for each patient, based on a variety of physical and metabolic factors.

It would have been prima facie obvious to a person of ordinary skill in the art at the time the invention was made to use a pyrimidine nucleoside such as tunicamycin to inhibit angiogenesis.

A person of ordinary skill in the art would have been motivated to use a pyrimidine nucleoside such as tunicamycin given the prior art's recognition of tunicamycin as an

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inhibitor of the pathway leading to the angiogenic process of capillary endothelial cell proliferation.

Applicant argues that Banerjee does not teach or suggest inhibiting angiogenesis using a nucleoside comprising glucosamine or a nucleoside comprising a pyrimidine nucleoside let alone tunicamycin and functional derivatives thereof; moreover that Banerjee is chiefly concerned with the evaluation of isoproterenol and it's mediation of protein glycosylation.

It should be noted that applicant has not responded to the specific citations in the reference showing Tunicamycin as an inhibitor of N-glycosylation; moreover, applicant would contend that because the aim of the study was not Tunicamycin, any teachings presented in the reference showing Tunicamycin as an inhibitor of N-glycosylation are not relevant. As cited in the previous office action, applicant should note that tunicamycin is a glucosamine containing nucleoside compound (emphasis added). Banerjee teaches that protein N-glycosylation and angiogenesis are indeed interlinked (p.293, ¶ 3); moreover, that Tunicamycin inhibited N-glycosylation in control cells by 64% and those treated with isoproterenol by 70% (p. 392, col.1-col.2). Thus Banerjee has recognized that Tunicamycin is a potent N-glycosylation inhibitor, as such given the teachings by Banerjee that angiogenesis is linked to N-glycosylation, one of skill in the art would have a reasonable expectation of success in the use of Tunicamycin to inhibit angiogenesis. Moreover, as cited supra, Tiganis et al., further supports the recognition in the prior art of the inhibition of N-glycosylation by tunicamycin and the disruption of vascular proliferation or angiogenesis.

The teachings of Tiganis verify that Tunicamycin is a potent inhibitor of glycoprotein synthesis in endothelial cells (see Discussion, p. 198) as equally purported by the teachings of Banerjee. As cited supra, Banerjee teaches that angiogenesis is a two part process, endothelial cell proliferation and differentiation. Clearly, one of skill in the art presented with the teachings of Tiganis on the ability of Tunicamycin to inhibit the proliferation of endothelial cells would find sufficient motivation in the use of Tunicamycin for the disruption or inhibition of angiogenesis.

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THIS ACTION IS MADE FINAL. Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the mailing date of this final action.

Howard V. Owens Patent Examiner Art Unit 1623

Samuel Barts

Primary Patent Examiner Technology Center 1600

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Howard Owens whose telephone number is (703) 306-4538 . The examiner can normally be reached on Mon.-Fri. from 8:30 a.m. to 5 p.m.

If attempts to reach the examiner by telephone are unsuccessful, the Supervisory Patent Examiner signing this action, James O. Wilson can be reached on (703) 308-4624. The fax phone number for this Group is (703) 308-4556.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the Group receptionist whose telephone number is (703) 308-1235.